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and also appears in each January and July issue.

Manuscripts—Ordinarily manuscripts should not exceed 3,000 words (less than 15 typed double-spaced pages). The original manuscript and two copies should be submitted typed double-spaced throughout. If photographic illustrations are desired, three sets of clear, unmounted black and white glossy prints should be supplied. For line drawing, original artwork or glossy prints are acceptable. Table should be typed on separate sheets of paper.

References—References should conform to the following order: name of author, title of article, name of periodical and volume with inclusive pages, month and year—in Lee GS: The heart rhythm following therapy with digitalis—In theory and practice. Arch Int Med 44:554-56. Dec 1942. They should be listed in the order in which they are cited in the text, numbered in sequence an punctuated as above.

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For Oral Use

DESCRIPTION: PROCARDIA (nifedipine) is an antianginal drug belonging to a new class of pharmacological agents, the calcium channel blockers. Nifedipine is 3,5-pyridinadicarboxylic acid, 1,4-dihydro-2,6-dimethyl-4-(2-nitrophenyl)-, dimethyl ester, C₁₇H₁₈N₂O_e, and has the struc-

Nifedipine is a yellow crystalline substance, practically insoluble in water but soluble in ethanol, it has a molecular weight of 346.3. PROCARDIA CAPSULES are formulated as soft getatin capsules for oral administration each containing 10 mg nifedipine.

CLINICAL PHARMACOLOGY: PROCARDIA (nifedipine) is a calcium ion linflux inhibitor (slow channel blocker or calcium ion antagonist) and inhibits the transmembrane influx of calcium ions into cardiac muscle and smooth muscle. The contractile processes of cardiac muscle and vascular smooth muscle are dependent upon the movement of extracellular calcium ions into these cells through specific ion channels. PROCARDIA selectively inhibits calcium ion influx across the cell membrane of cardiac muscle and vascular smooth muscle without changing serum calcium

cells through specific ion channels. PROCARDIA selectively inhibits calcium ion influx across the cell membrane of cardiac muscle and vascular amooth muscle without changing serum calcium concentrations.

Mischanism of Action: The precise means by which this inhibition relieves angina has not been fully determined, but includes at least the following two mechanisms:

1) Relaxation and prevention of coronary artery apasan: PROCARDIA dilates the main coronary arteries and coronary arterioles, both in normal and ischemic regions, and is a potent inhibitor of coronary artery spasm, whether spontaneous or ergonovine-induced. This property increases myocardial oxygen detivary in patients with coronary artery spasm, and is responsible for the effectiveness of PROCARDIA in vascspastic (Prinzmetal's or variant) angina. Whether this effect plays any role in classical angina is not clear, but studies of exercise tolerance have not shown an increase in the maximum exercise rate-pressure product, a widely accopited measure of oxygen utilization. This suggests that, in general, relief of spasm or dilation of coronary arteries is not an important factor in classical angina.

2) Reduction of oxygen utilization: PROCARDIA regularly reduces arterial pressure at rest and at a given level of exercise by dilating peripheral arterioles and reducing the total peripheral registrance (afterload) against which the heart works. This unloading of the heart reduces myocardial energy consumption and oxygen requirements and probably accounts for the effectiveness of PROCARDIA in chronic stable angina.

Pharmacokinetics and Metabolism: PROCARDIA is rapidly and fully absorbed after oral administration. The drug is detectable in serum 10 minutes after oral administration, and peak blood levels occur in approximately 30 minutes. It is highly bound by serum proteins. PROCARDIA and metabolites are eliminated via the kidneys. The half-life of nifediphre in plasma is approximately two hours. There is no information on the effects of renal or hepatic

is approximately two hours. There is no information on the effects of renal or hepatic impairment on excretion or metabolism of PROCARDIA.

Hemodynamics: Like other slow channel blockers, PROCARDIA exents a negative inotropic effect on isolated myocardial tissue. This is rarely, if ever, seen in intact animals or man, probably because of reflex responses to its vasoditating effects. In man, PROCARDIA causes decreased peripheral vascular resistance and a fall in systolic and diastolic pressure, usually modest (5–10 mm Hg systolic), but sometimes larger. There is usually a small increase in heart rate, a reflex response to vasoditation. Measurements of cardiac function in patients with normal ventricular function have generally found a small increase in cardiac index without major effects on ejection fraction, left ventricular end diastolic pressure. (LVEPP) or volume (LVEDV). In patients with impaired ventricular function, most acute studies have shown some increase in ejection fraction and reduction in fert ventricular end diastolic pressure.

Electrophysiologic Effects: Although, like other members of its class, PROCARDIA decreases sinoatrial node function and atroventricular conduction in isolated myocardial preparations, such effects have not been seen in studies in intact animals or in man. In formal electrophysiologic studies, predominantly in patients with normal conduction systems, PROCARDIA has had no tendency to prolong atrioventricular conduction, prolong sinus node recovery time, or slow sinus rate.

INDICATIONS AND USAGE: I. Vasospastic Angina: PROCARDIA (nifedipine) is indicated for the management of vasospastic angina confirmed by any of the following criteria: 1) classical patients who have had angiography, the presence of significant fixed obstructive disease is not incompatible with the diagnosis of vasospastic angina provided that the above criteria are satisfied. PROCARDIA may also be used where the clinical presentation suggests a possible vasospastic component but where vasospasm, or when a

blockers.

II. Chronic Stable Angine (Classical Effort-Associated Angine): PROCARDIA is indicated for the management of chronic stable angina (effort-associated angina) without evidence of vasospasm in patients who remain symptomatic despite adequate doses of beta blockers and/or organic nitrates or who cannot tolerate those agents.

In chronic stable angine (effort-associated angina) PROCARDIA has been effective in controlled trials of up to eight weeks duration in reducing angina frequency and increasing exercise tolerance, but confirmation of sustained effectiveness and evaluation of long-term safety in these

tolerance, but confirmation of sustained effectiveness and evaluation of long-term safety in these patients are incomplete.

Controlled studies in small numbers of patients suggest concomitant use of PROCARDIA and beta-blocking agents may be beneficial in patients with chronic stable angina, but available information is not sufficient to predict with confidence the effects of concurrent treatment in patients with compromised left ventricular function or cardiac conduction abnormalities. When introducing such concomitant therapy, care must be taken to monitor blood pressure since severe hypotension can occur from the combined effects of the drugs. See Warnings.

CONTRAINDICATIONS: Known hypersensitivity reaction to PROCARDIA with the description of the processive and poorly tolerated hypotension. These responses have usually occurred during initial thration or at the time of subsequent upward dosage adjustment, and may be more likely in patients on concomitant beta blockers.

Increased Angins/Beta Blocker Withdrawat: Occasional patients have developed well doc-

of subsequent upward dosage adjustment, and may be more likely in patients on concomitant beta blockers.

Increased Angins/Beta Blocker Withdrawrat: Occasional patients have developed well documented increased frequency, duration or severity of angina on starting PROCARDIA or at the time of dosage increased. The mechanism of this response is not established but could result from decreased coronary perfusion associated with decreased disabilic pressure with increased heart rate, or from increased demand resulting from increased heart rate alone.

Patients recently withdrawn from beta blockers may develop a withdrawal syndrome with increased angina, probably related to increased sensitivity to catecholamines. Initiation of PROCARDIA treatment will not prevent this occurrence and might be expected to exacerbate it by provoking reflex catecholamine release. There have been occasional reports of increased angina in a setting of bota blocker withdrawal and PROCARDIA initiation. It is important to taper beta blocker provided to the provided provided provided provided to the provided provided provided provided provided to the provided provided

diuretic therapy. With patients whose angina is complicated by congestive heart failute, should be taken to differentiate this peripheral edema from the effects of increasing left venue.

should be taken to differentiate this peripheral edema from the effects of increasing left ventila dysfunction.

Drug interactions: Beta-advenergic blocking agents: See Indications and Warnings. Experis in over 1400 patients in a non-comparative clinical frial has shown that concomitant administration of PROCARDIA and beta-blocking agents is usually well tolerated, but there have been on sional literature reports suggesting that the combination may increase the likelihood of comparative tailure, severe hypotension or exacerbation of angina.

Long-acting nitrates: PROCARDIA may be safely co-administered with nitrates, but there have been no controlled studies to evaluate the antianginal effectiveness of this combination.

Carcinogenesis, mutagenesis, impairment of fertility: Nielecipine was administered orally for for two years and was not shown to be carcinogenic. When given to rats prior to main nifedipine caused reduced fertility at a dose approximately 30 times the maximum recomment human dose. In vivo mutagenicity studies were negative.

Pregnancy: Pregnancy category C. Nifedipine has been shown to be teratogenic in rabs and piven in doses 30 times the maximum recommended human dose. Nifedipine was embrate (increased letal resorptions, decreased fetal weight, increased stunted forms, increased in maximum recommended human dose resulted in small placentas and underdeveloped chorionic will maximum recommended human dose in pregnant monkeys, doses 2/3 and twice the maxim resormmended human dose resulted in small placentas and underdeveloped chorionic and maximum pregnancy only if the potential benefit justifies the potential insk to the light attacts, docses three times the maximum human dose and higher caused prolongation or pregnant reactions were reported spontaneously, adverse effects were frequent but generally not see and rarely required discontinuation of therapy or dosage adjustment. Most were expected to and rarely required discontinuation of therapy or dosage adjustment. Most were expected to provide

Dizziness, light-headedness, giddiness Flushing, heat sensation Headache 20 10 Nausea, heartburn Muscle cramps, tremor Peripheral edema Nervousness, mood changes Palnitation Dyspnea, cough, wheezing Nasal congestion, sore throat

Nasal congestion, sore throat

There is also a large uncontrolled experience in over 2100 patients in the United States. Mo
the patients had vasospastic or resistant angina pectoris, and about half had concomitant in
ment with beta-adrenergic blocking agents. The most common adverse events were the ae
neas seen in the controlled trials, with dizziness or light-headedness, peripheral edema, nas
weakness, headache and flushing each occurring in about 10% of patients, tragatient hypoten
in about 5%, patientstion in about 2% and syncope in about 5.5%, Syncopal epidodes did note
with reduction in the dose of PROCARDIA or concomitant antianginal medication. Very rangitroduction of PROCARDIA therapy was associated with an increase in anginal pain, possibly
in associated hypotension.

with reduction in the dose of PROCARDIA or concomitant antianginal measurement reduction of PROCARDIA therapy was associated with an increase in anginal pain, possibly to associated hypotension.

Several of these side effects appear to be dose related. Peripheral edema occurred in all one in 25 patients at doses less than 60 mg per day and in about one patient in eight at 120 mg/day or more. Transient hypotension, generally of mild to moderate severity and seldom required day or more. Transient hypotension, generally of mild to moderate severity and seldom required sociations at 120 mg/per day or more.

In addition, 2% or fewer of patients reported the following: Respiratory: Nas-d and the congestion, shortness of breath. Gastrointestinal: Diarrhea, constipation, cramps fistulent functions, sleep disturbances, blurred vision, difficulties in balance. Other: Dermatitis, pruritig, ticaria, fever, sweating, chills, sexual difficulties.

In addition, more serious adverse events were observed, not readily distinguishable from natural history of the disease in these patients. It remains possible, however, that some or man these events were drug related. Myocardial infarction occurred in about 4% of patients a congestive heart failure or pulmonary edema in about 2%. Ventricular arrhythmias or conductive theart failure or pulmonary edema in about 2%. Ventricular arrhythmias or conductive for adverse experiences was not different from that of the edigroup of PROCARDIA treated patients (see Precautions).

In a subgroup of patients with a diagnosis of congestive heart failure as well as angina, deness or light-headedness, peripheral adema, headache or flushing each occurred in one ineit patients. Hypotension occurred in about one in 20 patients. Syncope occurred in approximance patients. Hypotension occurred in about one in a position of 250. Myocardial infarction or symptoms of congestive heart failure each occur in about one patient in 250. Myocardial infarction or symptoms of congestive heart failure each occur in abou

in about one patient in 15. Atrial or ventricular dysrhythmias each occurred in about one patien 150.

Laboratory tests: Rare, mild to moderate, transient elevations of enzymes such as alial phosphatase. CK, LDH, SGOT, and SGPT have been noted, and a single incident of significant displants are saved in a patient with a history of bladder disease after about elevere months of nitedipine therapy. The relationship PROCARDIA therapy is uncertain. These laboratory abnormalities have already been associal with clinical symptoms. Cholestasis, possibly due to PROCARDIA therapy, has been report twice in the extensive world literature.

OVERDOSAGE: Although there is no well documented expenence with PROCARDIA overdings, available data suggest that gross overdosage could result in excessive peripheral vised iton with subsequent marked and probably prolonged systemic hypotension. Clinically signification with subsequent marked and probably prolonged systemic hypotension. Clinically signification in the straing document marked and probably prolonged systemic hypotension. Clinically signification of the prolonged cardiac and respiratory function, elevation of extremities, and attention to circuit fluid volume and urine output. A vascoonstrictor (such as nonepinephrine) may be helpful in storing vascular tone and blood pressure, provided that there is no contraindication to its at Clearance of PROCARDIA would be expected to be prolonged in patients with impaired function. Since PROCARDIA is highly protein-bound, dialysis is not likely to be of benefit DOSAGE AND ADMINISTRATION: The doeage of PROCARDIA needed to suppress quant and that can be tolerated by the patient must be established by the tration. Excess quant and that can be tolerated by the patient must be established by the tration. Excess quant and that can be tolerated by the patient must be established by the patient with engage are rarely necessary. More than 180 mg per day is not recommended.

In most cases, PROCARDIA is tration should proceed over a 7–14 day per

cian can assess the response to each dose level and monitor the blood pressure before profiling to higher doses.

If symptoms so warrant, titration may proceed more rapidly provided that the patient is sessed frequently. Based on the patient's physical activity level, attack frequency, and subling introglycerin consumption, the dose of PROCARDIA may be increased from 10 mg t.i.d. to 20 t.l.d. and then to 30 mg t.i.d. over a three-day period.

In hospitalized patients under close observation, the dose may be increased in 10 mg ments over four to six-hour periods as required to control pain and arrhythmias due to ischemisingle dose should rarely exceed 30 mg.

No "rebound effect" has been observed upon discontinuation of PROCARDIA. However, if continuation of PROCARDIA is necessary, sound clinical practice suggests that the dose should be decreased gradually with close physician supervision.

Co-Administration with Other Antisinglina Druges: Sublingual nitroglycerin may be taked required for the control of acute manifestations of angine, particularly during PROCARDIA ton. See Precautions, Drug Interactions for information on co-administration of PROCARDIA with beta blockers or long-acting nitrates.

son. See Precurons, Drug interactions for information of co-administration of with beta blockers or long-acting intrates.

HOW SUPPLIED: Each orange, soft gelatin PROCARDIA Capsule contains 10 mg of niled PROCARDIA Capsules are supplied in amber glass bottles of 100 capsules (NDC 0069-2600 The capsules should be protected from light and moisture and stored at controlled temperature 59° to 77°F (15° to 25°C) in the manufacturer's original container.

Issued January 1982

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ESTHESIOLOGY

rch 11-14-Obstetrical Anesthesia: The State of the Art. 982. UCSF at Golden Gateway Holiday Inn, San Francisco. hursday-Sunday, 13 hrs. \$275.

rch 14-18-International Anesthesia Research Society-56th congress. Hilton Hotel, San Francisco. Sunday-Thursday. contact: International Anesthesia Research Soc., 3645 Warensville Center Rd., Cleveland, OH 44122. (216) 295-1124.

rch 17-21-American Society of Regional Anesthesia. Del Monte Hyatt, Monterey. Wednesday-Sunday. 16 hrs. \$175. contact: Gale Thompson, MD, ASRA, P.O. Box 11083, lichmond, VA 23230. (804) 358-0305.

rch 21-26—Anesthesiology 1982—8th Annual Winter Meetig. Anesthesiology Educational Foundation at Sierra Nevada nn, Mammoth Lakes. Sunday-Friday. Contact: AEF, P.O. ox 24230, Los Angeles 90024.

il 3-10-10th Obstetric Anesthesia Conference. Dept. of nesthesiology, Ohio State Univ. College of Medicine, at heraton Waikiki, Oahu, Hawaii. One week. 25 hrs. Contact: Pept. of Anesthesiology, c/o Bradford Travel, 1200 W. tenderson Rd., Columbus, OH 43220. (614) 457-9990 or \$14) 421-8487 (OSU).

il 26-30—6th Annual Hawaiian Seminar on Clinical Anesesiology. Calif. Society of Anesthesiologists at Maui Surf lotel, Maui. Monday-Friday. 20 hrs. Contact: Educational rograms Div., CSA, 100 E. Ellsworth Ave., Suite 806, San lateo 94401. (415) 348-1407.

22-23—Hemodynamic Monitoring/Pulmonary Artery atheterization—Hands-on Workshops. Calif. Soc. of Anesesiologists and Stanford at Stanford, Saturday-Sunday. 15 s. \$275, CSA members; \$325, nonmembers. Contact: Edutional Prog. Div., CSA, 100 S. Ellsworth Ave., Suite 806, in Mateo 94401. (415) 348-1407.

10-13—California Society of Anesthesiologists—Annual leeting. Town and Country, San Diego. Thursday-Sunday. ontact: Educational Prog. Div., CSA, 100 S. Ellsworth ve., Suite 806, San Mateo 94401. (415) 348-1407.

24-26-Practical Closed Circuit Anesthesia II. Pacific ledical Center at Hyatt at Union Square, San Francisco. hursday-Saturday. 17 hrs. \$300. Contact: Office of Cont. d., PMC, P.O. Box 7999, San Francisco 94120. (415) 563-21, ext. 2761.

ust 21-22—Hemodynamic Monitoring/Pulmonary Artery atheterization-Hands-on Workshops. Calif. Soc. of Anesesiologists and Stanford at Stanford. Saturday-Sunday. 15 8. \$275, CSA members; \$325, nonmembers. Contact: Edutional Prog. Div., CSA, 100 S. Ellsworth Ave., Suite 806, n Mateo 94401. (415) 348-1407.

CANCER

February 17-Advances in Cancer Detection. Hospital of the Good Samaritan at Biltmore Hotel, Los Angeles. Wednesday. 7 hrs. Contact: Bonnie VanWaardenburg, Cancer Symp., Hosp. of the Good Samaritan, 616 S. Witmer St., Los Angeles 90017. (213) 977-2345.

February 24-26—Current Approaches to Radiation Oncology. Biology and Physics. UCSF. Wednesday-Friday.

February 27-28-Breast Cancer: Conservation Surgery and Radiation Therapy Now-17th Annual San Francisco Cancer Symposium. West Coast Cancer Fndn.; Am. Cancer Soc., Calif. Div.; St. Mary's Hosp, and Med. Ctr. at Sheraton Fisherman's Wharf, San Francisco. Saturday-Sunday. 12 hrs. \$115 before Jan. 1; \$140, thereafter. Contact: West Coast Cancer Fndn., 50 Francisco St., Suite 200, San Francisco 94133. (415) 981-4590.

March 25-27-Western States Conference on Cancer Rehabilitation: Psychosocial, Physical and Economic Interventions. No. Calif. Cancer Prog.; Am. Cancer Soc., Calif. Div.; UCLA Jonsson Comprehensive Cancer Ctr.; L.A. Co.-USC Comprehensive Cancer Ctr.; Community Cancer Control-

(Continued on Page 25)

KEY TO ABBREVIATIONS

CMA: California Medical Association

Contact: Continuing Medical Education, California Medical Association, 731 Market Street, San Francisco 94103. (415) 777-2000.

DREW: Charles R. Drew Postgraduate Medical School Contact: Duane Dillman, PhD, Director, Office of Continuing Education, Charles R. Drew Postgraduate Medical School, 1621 East 120th Street, Los Angeles 90059. (213) 603-3091.

LLU: Loma Linda University Contact: Varier J. Johns, Jr., MD, Associate Dean for Continuing Medical Education, Loma Linda University School of Medicine, Loma Linda 92354. (714) 796-7311. Ext. 2400.

Pacific Medical Center PMC: Contact: Continuing Education, Pacific Medical Center, P.O. Box 7999, San Francisco 94120. (415) 563-4321, ext. 2761.

STAN: Stanford University
Contact: Edward Rubenstein, MD, Associate Dean for
Postgraduate Education, Stanford University School of
Medicine, 300 Pasteur Drive, Stanford 94305. (415)
497-5594.

University of California, Davis UCD: Contact: Neil C. Andrews, MD, Chairman, Office of Continuing Medical Education, University of California, Davis, School of Medicine, Davis 95616. (916) 752-0328.

University of California, Irvine UCI: Contact: E. R. Jennings, MD, Assistant Dean, University of California, Irvine, California College of Medicine, c/o UCI/CME Program, Memorial Hospital Medical Center 2801 Atlantic Ave., Long Beach 90801. (213) 595-3837.

UCLA: University of California, Los Angeles
Contact: Martin D. Shickman, MD. Director, Continuing Education in Medicine and the Health Sciences, P.O.
Box 24902, UCLA, Los Angeles 90024. (213) 825-7241.

UCSD: University of California, San Diego
Contact: David Allan, MD, Associate Dean for Continuing Medical Education, University of California, San Diego, School of Medicine (M-017), La Jolia 92093. (714) 452-3708.

UCSF: University of California, San Francisco
Contact: Malcolm S. M. Watts, MD. Associate Dean
and Director, Extended Programs in Medical Education,
School of Medicine, University of California, San Francisco 94143. (415) 666-4251.

University of Southern California USC: Contact: Phil R. Manning, MD, Associate Dean, Postgraduate Division, University of Southern California School of Medicine, 2025 Zonal Avenue, Los Angeles 90033. (213) 224-7051.

- L.A.; S.F. Regional Cancer Findin at Fairmont Hotel, San Francisco. Thursday-Saturday. Contact: Carrie Ewing, No. Calif. Cancer Prog., P.O. Box 10144, Palo Alto 94303. (415) 497-7431.
- April 2-7—Hematology/Oncology Update. See Medicine, April 2-7.
- April 3—Oncology Update 1982. Northridge Hospital Medical Center at Biltmore Hotel, Los Angeles. Saturday, 8 hrs. \$125. Contact: Sandra Rozzen, Med. Ed. Dept., NHMC, 18300 Roscoe Blvd., Northridge 91328. (213) 885-5311.
- pril 22-24—Bladder Cancer. American Urological Assn. at Biltmore Hotel, Los Angeles. Thursday-Saturday. 16 hrs. Contact: Alice Henderson, AUA Office of Education, P.O. Box 25147, Houston, TX 77005.

MERGENCY MEDICINE

- tioner. Calif. Chap., American College of Emergency Physicians, at St. Mary Medical Center, Long Beach. Saturday. 8 hrs. \$75. ACEP members; \$95, nonmembers. Contact: CAL/ACEP, 8639 Lincoln Blvd., Suite 240, Los Angeles 90045. (213) 776-1820.
- ebruary 28-March 5—Emergency Medicine. UCD at Hyatt, Lake Tahoe. Sunday-Friday.
- arch 7-10—Sports Injuries Update. UCD at Hyatt, Lake Tahoe. Sunday-Wednesday.
- March 14-19—Third Annual Mammoth Mountain Emergency Medicine Ski Conference. UCI at Mammoth Lakes. Sunday-Friday. 25 hrs. \$290. Contact: Medical Conferences, Inc., P.O. Box 52-B, Newport Beach 92662. (714) 540-9876.
- farch 15-19—Postgraduate Institute for Emergency Room Physicians, Symposium I. UCSD at Islandia Hotel, San Diego. Monday-Friday. 36 hrs. \$425.
- pril 10-17—Pediatric Emergencies, UCSD at Kauai, Hawaii. One week. 21 hrs. \$325.
- pril 23-25—Advanced Cardiac Life Support Workshops. Emergency Medical Systems, Inc., at Ralph K. Davies Medical Center, San Francisco. Friday-Sunday. Contact: EMS, Inc., 2310 Mason St., San Francisco 94133.
- pril 24-May 2—Emergency Medicine. USC at Hawaii. One week. 25 hrs. \$425.
- by 10-14—Postgraduate Institute for Emergency Room Physicians, Symposium I. UCSD. Monday-Friday. 36 hrs. \$425.
- te 7-11—Postgraduate Institute for Emergency Room Phyicians—Symposium III. UCSD at Islandia Hyatt, San Diego. Monday-Friday. 36 hrs. \$425.
- te 28-July 2—Topics in Emergency Medicine. UCSF. Monday-Friday. 35 hrs. \$350.
- 23-25—Advanced Cardiac Life Support Workshops. Emerency Medical Systems, Inc., at Ralph K. Davies Medical center, San Francisco. Friday-Sunday. Contact: EMS, Inc., 310 Mason St., San Francisco 94133.
- gust 9-13—Postgraduate Institute for Emergency Room Physicians—Symposium II. UCSD at Islandia Hyatt, San Piego. Monday-Friday. 36 hrs.

DICINE

- tuary 16-19—Clinical Investigation. See Of Interest to All, bruary 16-19.
- triary 17-19—Recent Advances in Noninvasive Cardiac taging: Applications in the Diagnosis and Management of themic Heart Disease. American College of Cardiology at antury Plaza Hotel, Los Angeles. Wednesday-Friday. Control Registration Secretary, Extramural Prog. Dept., ACC, 11 Old Georgetown Rd., Bethesda, MD 20014. (301) 897-00.
- ary 18-20—Controversies and Problem Solving With Encopy—An Update. So. Calif. Society for Gastrointestinal

- Endoscopy at Beverly Hilton Hotel, Beverly Hills. Thursday-Saturday. 18 hrs. Contact: Dr. Albert Dreskin, 13652 Cantara St., Panorama City 91402.
- February 20-27—Cardiology. Univ. of Washington School of Medicine at Wailea Beach, Maui. One week. Contact: Univ. of WA Sch. of Med., Div. of CME, SC-50, Seattle 98195. (206) 543-1050.
- February 21-23—Recent Advances in Clinical Endocrinology. UCSF. Sunday-Tuesday.
- February 24-25—American College of Physicians—Regional Meeting. Hilton, Beverly Hills. Wednesday-Thursday. Contact: Mitchel D. Covel, MD, 9730 Wilshire Blvd., Beverly Hills 90212.
- February 25-28—Federation of the Western Societies of Neurological Science. Hotel del Coronado, Coronado. Thursday-Sunday. 15 hrs. Contact: Renee Kaseff, 525 Hurlingham Ave., San Mateo 94402. (415) 344-9082.
- February 26—A Day in Nephrology. Sutter Community Hospitals at Holidome Convention Center, Sacramento. Friday. 7 hrs. Contact: Barbara Davis, (916) 446-6667.
- February 26-27—American College of Physicians—Regional Meeting. Doubletree Inn. Monterey. Friday-Saturday. Contact: Jerry P. Lewis, MD, 618 Francisco Place, Davis 95616.
- February 26-28—Arrhythmias and Cardiac Ischemia: Diagnosis and Management. International Medical Education Corp. at Aladdin Hotel, Las Vegas. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- February 27-28—Immunology. STAN. Saturday-Sunday. 12 hrs. \$225.
- February 27-28—Basic Neuroscience. STAN. Saturday-Sunday. 12 hrs. \$225.
- February 27-28—Asthma: Impact and Strategies for Change. UCLA. Saturday-Sunday. 14 hrs. \$85.
- March 3-6—Third World Congress for Bronchology. World Assn. for Bronchology and American College of Chest Physicians at Town and Country, San Diego. Wednesday-Saturday. \$250, WAB and ACCP members; \$300, nonmembers. Contact: ACCP, P.O. Box 93826, Chicago 60670. (312) 698-2200.
- March 3-6—Immunology and the Clinical Practice of Allergy—6th Annual Course. UCSD at Hilton Hotel, San Diego. Wednesday-Saturday. 20 hrs.
- March 3-6—Third World Congress for Bronchology. American College of Chest Physicians and World Assn. for Bronchology at Town and Country Hotel, San Diego. Wednesday-Saturday. \$250, ACCP and WAB members; \$300, nonmembers. Contact: ACCP, P.O. Box 93826, Chicago 60670. (312) 698-2200.
- March 5—Diabetes. UCD at Red Lion Inn, Sacramento. Friday.
- March 7-10—Basic Cardiology for the Practicing Physician. UCSF at Lake Tahoe. Sunday-Wednesday.
- March 10—Hypothyroidism—Unmasked. LLU. Wednesday. 4 hrs. \$30.
- March 12-13—Exercise Stress Testing: Today and Tomorrow. American College of Cardiology at Memorial Hospital Medical Center, Long Beach. Friday-Saturday. Contact: Registration Secretary, Extramural Prog. Dept., ACC, 9111 Old Georgetown Rd., Bethesda, MD 20014. (301) 897-5400.
- March 12-13—Clinical Management of Coronary Disease and Exercise Testing. International Medical Education Corp. at Sheraton at the Wharf, San Francisco. Friday-Saturday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- March 12-14—Western Institute on Epilepsy. UCLA at Marriott, Los Angeles. Friday-Sunday. \$60.
- March 13-20—Reproductive Endocrinology. USC at Maui. One week. \$395.

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- arch 19-20—Non-Invasive Cardiology Conference. Clairemont Community Hospital at Sheraton Harbor Island Hotel, San Diego. Friday-Saturday. Contact: Nomi Feldman, 3770 Transy, San Diego 92121. (714) 453-6222.
- arch 21-24—Dermatology Consultants' Course. UCSF at Cosemite. Sunday-Wednesday.
- farch 22-April 2—30th Annual Review of Internal Medicine.
- ment. International Medical Education Corp. at Sahara Hotel, Las Vegas. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- March 29-April 2—Management of Coronary Heart Disease: Medical, Surgical, Exercise. UCSD at Hilton, San Diego. Monday-Friday. 40 hrs.
- pril-June—Allergy/Immunology for the Practicing Internist.
- pril 1-3—Asthma Update—1982. Memorial Hospital Medical Center-UCI Center for Health Education, Long Beach. Thursday-Saturday. 19 hrs. Contact: Asst. Dir., Ctr. for Health Education, 2801 Atlantic Ave., Long Beach 90801. (213) 595-3811.
- April 1-4—Principles in the Practice of Allergic Diseases in Adults and Children. UCD at Del Monte Hyatt, Monterey. Thursday-Sunday.
- April 2-7—Hematology/Oncology Update. UCD at Ahwahnee Hotel, Yosemite. Friday-Wednesday.
- American Heart Assn.-Greater Los Angeles Affiliate, San Gabriel Valley Division, at Huntington Sheraton, Pasadena. Saturday. 8 hrs. \$55, members; \$70, nonmembers. Contact: Gary L. Kossky, AHA, 929 N. Grand Ave., Covina 91724. (213) 339-7382 or (213) 686-2506.
- April 14—Cardiology Update 1982. N. T. Enloe Memorial Hospital, Chico. Wednesday. 6-7 hrs. \$35. Contact: Med. Staff Office, Enloe Hosp., W. 5th Ave. and The Esplanade, Chico 95926. (916) 891-7375.
- April 16—Injectable Collagen—Applications and Technique. STAN. Friday. \$100. Contact: Paul H. Jacobs, MD, Dept. of Dermatology, Stanford Univ. Sch. of Med., Stanford 94305. (415) 497-6101.
- oril 16-17—Recent Advances in Renal, Metabolic and Endocrine Disorders. St. Francis Medical Center at Bonaventure Hotel, Los Angeles. Friday-Saturday. \$115. Contact: Allan M. Hoffman, EdD, Dir., Med. Ed., St. Francis Med. Ctr., 3630 E. Imperial Highway, Lynwood 90262. (213) 603-6171.
- pril 16-18—Arrhythmias and Cardiac Ischemia: Diagnosis and Management. International Medical Education Corp. at Sheraton, Anaheim. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- 22-23—Infectious Disease Symposium. Epidemiology Assn. of Greater Sacramento and Mercy Hospital at Red Sion Inn, Sacramento. Thursday-Friday. 14 hrs. Contact: Office of Cont. Med. Ed., Mercy General Hospital, 4001 J., Sacramento 95819. (916) 453-4545.
- 25-28—Dermatology. UCD at Ahwahnee Hotel, Yomite. One week.
- 28—Thyroid '82. Hospital of the Good Samaritan at onaventure Hotel, Los Angeles. Wednesday. 7 hrs. Contist: Bonnie VanWaardenburg, Coordinator of Symposia, of the Good Samaritan, 616 S. Witmer St., Los ageles 90017. (213) 977-2345.
- 6-7—Advances in Infectious Diseases, UCSF. Thursday-iday, 14 hrs. \$170.
- 7-8—Common Nervous System Pathology Diagnosis and Patment. Memorial Hospital Medical Center-UCI Center Health Education, Long Beach. Friday-Saturday. 14 hrs.

- Contact: Assi. Dir., Ctr. for Health Ed., 2801 Atlantic Ave., Long Beach 90801. (213) 595-3811.
- May 7-9—Ambulatory Electrocardiography: Clinical Applications, Methodology and Interpretation. International Medical Education Corp. at MGM Grand, Las Vegas. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- May 15-18—American Thoracic Society—Annual Scientific Meeting. Bonaventure Hotel, Los Angeles. Saturday-Tuesday. 30 hrs. \$35, members; \$100, nonmembers. Contact: S. R. Iannotta, Exec. Dir., ATS, 1740 Broadway, New York 10019. (212) 245-8000.
- May 17-21—Advances in Internal Medicine. UCSF. Monday-Friday. 40 hrs. \$375.
- May 21-22—EKG Interpretation and Arrhythmia Management. International Medical Education Corp. at Sheraton at the Wharf, San Francisco. Friday-Saturday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- May 27-30—4th Annual Refresher Course in Allergy and Asthma. UCD at Marriott's Great America, Santa Clara. Thursday-Sunday.
- June 7-9—Clinical Echocardiography—1982: M-Mode, 2-Dimensional and Pulse Doppler Techniques. American College of Cardiology at Sheraton Harbor Island Hotel, San Diego. Monday-Wednesday. Contact: Extramural Prog. Dept., ACC, 9111 Old Georgetown Rd., Bethesda, MD 20014.
- June 9—Bleeding Disorders. LLU. Wednesday. 4 hrs. \$30.
- June 11-13—Arrhythmias and Cardiac Ischemia: Diagnosis and Management. International Medical Education Corp. at Aladdin Hotel, Las Vegas. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- June 25-27—Clinical Management of Coronary Disease and Exercise Testing. International Medical Education Corp. at Newporter Inn, Newport Beach. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- July 16-17—Arrhythmias and Cardiac Ischemia: Diagnosis and Management. International Medical Education Corp. at Sheraton at the Wharf, San Francisco. Friday-Saturday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- July 17-24—Cardiovascular Medicine and Surgery: An Advanced Course. STAN at Mauna Kea Beach Hotel, Kamuela, Hawaii. One week. 21 hrs. \$325.
- August 6-8—EKG Interpretation and Arrhythmia Management. International Medical Education Corp. at Clouds-Cal Neva Resort, Lake Tahoe. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- August 13-15—Clinical Management of Coronary Disease and Exercise Testing. International Medical Education Corp. at Doubletree Inn, Monterey. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- August 20-22—Ambulatory Electrocardiography: Clinical Applications, Methodology and Interpretation. International Medical Education Corp. at Hyatt, Anaheim. Friday-Sunday. 13 hrs. \$245. Contact: Stephen E. Mattingly, IMEC, 64 Inverness Dr. E., Englewood, CO 80112. (800) 525-8651.
- August 30-September 4—Fifth Comprehensive Nephrology Review Course. UCLA at Marriott, Los Angeles. Monday-Saturday. 28½ hrs. \$595.

OBSTETRICS AND GYNECOLOGY

- March 13-20—Reproductive Endocrinology, Infertility and Contraception. See Of Interest To All, March 13-20.
- March 14-17—OB/GYN for the Practicing Physician. UCSF at Yosemite. Sunday-Wednesday.

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EPITOMES-PHYSICAL MEDICINE AND REHABILITATION

cations with anticoagulant drugs. A brief update on the diagnosis of pulmonary embolism is therefore included in this series of epitomes on physical medicine and rehabilitation.

Current practice in suspected PE may include initiation of anticoagulation on clinical criteria alone. However, physical signs and symptoms typically are nonspecific. More commonly, treatment is begun after arterial blood gas studies and ventilation-perfusion (V/Q) lung scanning. However, these approaches may result in misdiagnosis and inappropriate and hazardous treatment in many cases. A low arterial oxygen pressure (Po₂) may be present with a variety of acute or chronic lung diseases while a normal Po2 may occur in up to 15 percent of patients with documented PE. \dot{V}/\dot{Q} scanning has been regarded as an adequate screening test for PE in terms of sensitivity and specificity, although a recent report questions the validity of this concept. False-positive findings on \dot{V}/\dot{Q} scans are common, occurring in from 60 percent to 80 percent of cases when pulmonary angiography is used to confirm the presence of PE. False-negative findings of PE on V/Q scans, however, are rare. Thus, normal findings on \dot{V}/\dot{Q} scan virtually rule out the presence of PE if done between 24 and 72 hours from onset of symptoms.

The diagnosis of PE usually leads to such treatment as administration of anticoagulant drugs or vena cava ligation, both of which may have serious

complications. Thus, pulmonary angiography to assure that PE is present is recommended in situations where PE is suspected and the \dot{V}/\dot{Q} scan is positive for PE. These situations include cases in which patients have parenchymal lung disease or abnormalities in both ventilation and perfusion scans (because of the very high rate of false-positive scans), and cases in which patients are being considered for vena cava ligation or in whom anticoagulant drugs are contraindicated or who are at high risk for complications from anticoagulation therapy.

The importance of pulmonary angiography to confirm the presence of PE in patients at high risk for complications from anticoagulation therapy is just beginning to be realized. Carrying out potentially dangerous therapy in a patient who does not require it is hazardous. Heparin is the leading cause of adverse drug reactions in patients in hospital and anticoagulant drugs taken orally pose similar problems. Although pulmonary angiography is not a screening procedure, it is the procedure of choice for confirming the diagnosis of PE in high-risk patients. JAMES S. LIEBERMAN, MD

REFERENCES

ROBERT G. TAYLOR, MD

Giudice JC, Komansky HJ, Gordon R: Pulmonary thromboembolism—1. Current concepts in pathogenesis and diagnosis. Postgrad Med 67:64-71, 74-75, 77, May 1980

Robin ED: Overdiagnosis and overtreatment of pulmonary embolism: The emperor may have no clothes. Ann Intern Med 87:775-781, Dec 1977

Viamonte M Jr, Koolpe H, Janowitz W, et al: Pulmonary thromboembolism—Update. JAMA 243:2229-2234, Jun 6, 1980

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